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APPLICATION NO.	F	ILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/451,641	641 11/30/1999		Danchen Gao	C-3169/1/US	9327	
26648	7590	05/17/2005		EXAMINER		
		PORATION EPARTMENT	TRAN, SUSAN T			
POST OFFI				ART UNIT	PAPER NUMBER	
ST. LOUIS,	MO 630	006		1615		

DATE MAILED: 05/17/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application	on No.	Applicant(s)					
	09/451,64	1 1	GAO ET AL.					
Office Action Summary	Examiner	-	Art Unit					
	Susan T.	Tran	1615					
The MAILING DATE of this communication Period for Reply	appears on the	cover sheet with the c	orrespondence a	ddress				
A SHORTENED STATUTORY PERIOD FOR RE THE MAILING DATE OF THIS COMMUNICATIO - Extensions of time may be available under the provisions of 37 CFR after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a. - If NO period for reply is specified above, the maximum statutory per - Failure to reply within the set or extended period for reply will, by state Any reply received by the Office later than three months after the meanned patent term adjustment. See 37 CFR 1.704(b).	N. R 1.136(a). In no evo- reply within the state riod will apply and wi atute, cause the app	ent, however, may a reply be tim utory minimum of thirty (30) day: Ill expire SIX (6) MONTHS from lication to become ABANDONE!	nely filed s will be considered time the mailing date of this of					
Status								
1) Responsive to communication(s) filed on 2	4 February 20	<u>05</u> .						
2a)⊠ This action is FINAL . 2b)□ T	his action is n	on-final.						
3) Since this application is in condition for allo								
closed in accordance with the practice under	er <i>Ex parte</i> Qu	ayle, 1935 C.D. 11, 45	53 O.G. 213.					
Disposition of Claims	·							
4) Claim(s) <u>1-10,12-50,72-75,84 and 86-90</u> is/	are pending ir	the application.						
4a) Of the above claim(s) is/are without	drawn from co	nsideration.						
5) Claim(s) is/are allowed.								
6)⊠ Claim(s) <u>1, 2, 4-10,12-50,72-75,84 and 86-90</u> is/are rejected.								
7)⊠ Claim(s) <u>3</u> is/are objected to.								
8) Claim(s) are subject to restriction an	d/or election r	equirement.						
Application Papers								
9) The specification is objected to by the Exam	niner.							
10) The drawing(s) filed on is/are: a) a		Objected to by the 8	Examiner.					
Applicant may not request that any objection to								
Replacement drawing sheet(s) including the con		·		FR 1.121(d).				
11) The oath or declaration is objected to by the	•	=						
Priority under 35 U.S.C. § 119								
12) Acknowledgment is made of a claim for fore	ign priority un	der 35 U.S.C. § 119(a))-(d) or (f).					
a) ☐ All b) ☐ Some * c) ☐ None of:	•							
1. Certified copies of the priority docum								
2. Certified copies of the priority docum								
3. Copies of the certified copies of the p	=		ed in this National	l Stage				
application from the International Bur	•							
* See the attached detailed Office action for a	list of the certi	fied copies not receive	ed.					
Attachment(s)			(DTO 440)					
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948)		4) Interview Summary Paper No(s)/Mail Da						
3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/		5) Notice of Informal P		O-152)				
Paper No(s)/Mail Date <u>02/14/05</u> . 2/24/05		6) Other:		_				
U.S. Patent and Trademark Office PTOL-326 (Rev. 1-04) Office	e Action Summa	ry	Part of Paper No./Ma	il Date 050405				

DETAILED ACTION

Receipt is acknowledged of applicant's Petition to Withdraw from Issue, Request for Continued Examination, and Information Disclosure Statement filed 02/24/05.

Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after allowance or after an Office action under *Ex Parte Quayle*, 25 USPQ 74, 453 O.G. 213 (Comm'r Pat. 1935). Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, prosecution in this application has been reopened pursuant to 37 CFR 1.114. Applicant's submission filed on 02/24/05 has been entered.

Information Disclosure Statement

The information disclosure statement (IDS) submitted on 02/24/05 was filed after the mailing date of the allowance on 11/21/03. The submission is in compliance with the provisions of 37 CFR 1.97. Accordingly, the information disclosure statement is being considered by the examiner.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Claims 1, 2, 4-10, 12-50, 72-75, 84 and 86-90 are rejected under 35 U.S.C. 103(a) as being unpatentable over AAPS Annual Meeting Contributed Papers Abstracts (AAPS), in view of Black EP 0 863 134.

AAPS teaches a celecoxib (Cox-2 inhibitor) formulation that exhibits a C_{max} values of 1527 and 1077 ng/mL, and a T_{max} of 1.9 hours (see page D32).

AAPS does not teach the use of excipients in the formulation. However, the use of excipients in oral formulations is well known in pharmaceutical art. To be more specific, Black teaches a compound useful as a Cox-2 inhibitor for pain relief, fever and inflammation of a variety symptoms disclosed on page 3, lines 29-36. The compound can be administered orally in the form of tablets, troches, lozenges, or capsules (page 4, lines 1-12). The tablets comprising active ingredient in admixture with excipients, e.g., diluents, disintegrants, binding agents, wetting agents, and surfactant (page 4, lines 15-38). The active agent present in an amount of 10 to 250 mg, and carrier material may vary from about 5 to about 95% (page 5, lines 39-58). The dosage can be administered once or twice a day, and will provide effective T_{1/2} over a 24 hours period (page 5, lines 22-27). Example 2 discloses the amount of excipients use in a tablet. Thus, it would have been obvious for one of ordinary skill in the art to prepare the formulation of AAPS using excipient in view of the teachings of Black to obtain the claimed invention, because Black teaches oral dosage of Cox-2 inhibitor, and because

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AAPS teaches orally administering celecoxib in fine suspension and capsule forms having the claimed C_{max} and T_{max} values.

It is noted that AAPS does not expressly teach the particle size distribution, however, the burden is shifted to applicant to show that the formulation of AAPS does not have the claimed particle size distribution, as well as the detrimental effect and/or unexpected results over the particle size distribution, because AAPS teaches the oral formulation of celecoxib having the claimed C_{max} and T_{max} values.

Conclusion

All claims are drawn to the same invention claimed in the application prior to the entry of the submission under 37 CFR 1.114 and could have been finally rejected on the grounds and art of record in the next Office action if they had been entered in the application prior to entry under 37 CFR 1.114. Accordingly, **THIS ACTION IS MADE FINAL** even though it is a first action after the filing of a request for continued examination and the submission under 37 CFR 1.114. See MPEP § 706.07(b).

Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of

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the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Claims Allowable

Claim 3 is objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

Correspondence

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Susan T. Tran whose telephone number is (571) 272-0606. The examiner can normally be reached on M-R from 6:00 am to 4:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Thurman K. Page, can be reached at (571) 272-0602. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should

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you have questions on access to the Private PAIR system, contact the Electronic

Business Center (EBC) at 866-217-9197 (toll-free).

THURMAN K. PAGE SUPERVISORY PATENT EXAMINER TECHNOLOGY CENTER 1600

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